EXHIBIT 3

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14 15	Attorneys for Defendant NOVO NORDISK INC.	
16	UNITED STATES DISTRICT COURT	
17 18	FOR THE SOUTHERN DISTRICT OF CALIFORNIA	
19 20 21 22 23 24 25	In re: INCRETIN-BASED THERAPIES PRODUCTS LIABILITY LITIGATION As to All Related and Member Cases	Case No. 13-md-2452-AJB-MDD DEFENDANT NOVO NORDISK INC.'S SUPPLEMENTAL OBJECTIONS AND RESPONSES TO PLAINTIFFS' AMENDED SECOND SET OF INTERROGATORIES Judge: Hon. Anthony J. Battaglia Magistrate: Hon. Mitchell D. Dembin
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NOVO NORDISK INC.'S SUPPL. OBJ. AND RESP. TO PLS' AMENDED SECOND SET OF INTERROGATORIES

3:13-MD-02452-AJB-MDD

Defendant Novo Nordisk Inc. ("NNI"), for itself alone and for no other defendant, pursuant to Rules 26 and 33 of the Federal Rules of Civil Procedure, hereby supplements its objections and responses to Plaintiffs' Amended Second Set of Interrogatories filed with this Court on February 14, 2014, as follows:

GENERAL OBJECTIONS

- 1. NNI objects to these interrogatories to the extent they impose obligations beyond those imposed by the Federal Rules of Civil Procedure, any applicable Order of this Court, including, but not limited to, the Order governing the Production of Electronically Stored Information ("ESI Order"), the CMO Governing Limitations on Written Discovery, when entered, or this Court's Local Rules. *See* Order Governing the Production of Electronically Stored Information, *In re Incretin-Based Therapies Prods. Liab. Litig.*, No. 13-md-2452-AJB-MDD (S.D. Cal. Nov. 15, 2013), Doc. No. 187 ("ESI Order").
- 2. NNI objects to the voluminous nature of these interrogatories paired with the additional sets of interrogatories propounded on NNI, the total number of which exceeds that which is allowed under the Federal Rules of Civil Procedure. Nevertheless, following agreement by the parties, NNI agreed to respond to these interrogatories while reserving its rights to include substantive objections about which the parties will be prepared to meet and confer.
- 3. NNI objects to these interrogatories to the extent they seek information protected by the attorney-client, work-product, or any other applicable privilege or immunity from discovery. Any disclosure of information protected by any such privilege or other immunity shall be deemed inadvertent and shall not constitute a waiver of such privilege or other immunity.
- 4. NNI objects to these interrogatories, including subparts, to the extent they call for information that is not in the possession, custody, or control of NNI, or is a matter of public record or otherwise as accessible to Plaintiffs as to NNI.
 - 5. NNI objects to Plaintiffs' Definitions to the extent such definitions, as

incorporated into these interrogatories, renders an interrogatory vague, ambiguous, overly broad, unduly burdensome, and not reasonably calculated to lead to the discovery of admissible evidence. Specifically, NNI objects to Plaintiffs' definitions of "YOU," "YOUR," "YOURS," or "Defendants" as vague and ambiguous. NNI's responses to interrogatories herein construe terms "You," "Your," or "Defendant" to mean Novo Nordisk Inc. only.

- 6. NNI objects to these interrogatories to the extent they seek information from time periods that are irrelevant or inapplicable to Victoza®.
- 7. NNI objects to these interrogatories to the extent they seek information concerning products other than Victoza® (liraglutide).
- 8. NNI objects to these interrogatories to the extent they seek to function as document requests.
- 9. NNI objects to these interrogatories to the extent they call for the identification of all documents, individuals, information, or communication as well as any and/or every document, individual, piece of information, or communication when all relevant facts can be obtained from fewer than "all" documents or "any" document.
- 10. NNI objects to these interrogatories to the extent they seek information pertaining to injuries, alleged side effects, or adverse reactions not at issue in this litigation on the grounds that such interrogatories are not relevant, overly broad, and not reasonably calculated to lead to the discovery of admissible evidence.
- 11. NNI objects to these interrogatories to the extent they seek information related to foreign regulatory submissions, requirements, or activities, or the direction of foreign regulatory bodies, because it is neither relevant nor reasonably calculated to led to the discovery of admissible evidence and is unduly burdensome. Such information is subject to different regulatory and legal standards and requirements, and can be influenced by political, cultural, and social differences, including, but not limited to, differences in the practice of medicine.

- 12. NNI objects to these interrogatories to the extent they seek sales, marketing, or advertising information outside of the United States because it is neither relevant nor reasonably calculated to lead to the discovery of admissible evidence and is unduly burdensome to produce in this litigation.
- 13. NNI objects to these interrogatories to the extent the information sought is already in Plaintiffs' possession, custody or control, or are equally available to the Plaintiffs, on the grounds that such discovery requests are unreasonably cumulative and duplicative, and that the information may be obtained from a source that is more convenient, less burdensome, and less expensive.
- 14. NNI objects to these interrogatories to the extent they seek an analysis or summary of documents or information that is generally available to all parties. NNI objects further pursuant to Federal Rule of Civil Procedure 33(d) on the grounds that the burden of ascertaining such information is substantially the same for Plaintiffs as for NNI.
- 15. NNI objects to these interrogatories to the extent they seek confidential, proprietary, competitively sensitive, or trade secret information. To the extent NNI produces responsive and non-privileged information, any such information will be produced in accordance with the agreed-upon and Court-ordered Protective Order entered in this Litigation.
- 16. NNI objects to these interrogatories to the extent that they are duplicative of the deposition notices for testimony pursuant to Rule 30(b)(6) of the Federal Rules of Civil Procedure that Plaintiffs have also served on NNI in this action.
- 17. NNI objects to these interrogatories to the extent they request NNI to disclose the identity of any individual who allegedly experienced an adverse effect or who reported such an adverse experience on the ground that such a disclosure would violate the patients' or reporters' right to confidentiality under federal law.
 - 18. NNI's investigation into this matter is ongoing. Therefore, NNI may be

unable to provide full and complete responses to certain interrogatories. NNI will respond to these interrogatories as fully and completely as possible. NNI may supplement these responses as additional, responsive, relevant and non-privileged information becomes available.

- 19. By responding to these interrogatories, NNI does not concede the relevance, materiality, or admissibility of any of the documents sought herein for use as evidence in any hearing or trial. NNI's responses are made subject to, and without waiving, any objections as to relevance, materiality, or admissibility. NNI expressly reserves the right to object to further discovery on the subject matter of any of these requests.
- 20. NNI objects to these interrogatories to the extent they are outside the scope of general causation discovery and inapplicable discovery at this time pursuant to this Court's February 18, 2014 Order, which requires Plaintiffs to "narrow all discovery related requests to issues involving general causation." *See* Initial Case Management Scheduling Order Regarding General Causation, *In re Incretin-Based Therapies Prods. Liab. Litig.*, No. 13-md-2452-AJB-MDD (S.D. Cal. Feb. 18, 2014), Doc. No. 325 ("February 18 Order"). NNI will supplement its answers with information beyond general causation at the appropriate time, if necessary and applicable, per further Order of this Court.
- 21. The applicable foregoing General Objections are incorporated into each of the specific objections and responses that follow. Stating a specific objection or response shall not be construed as a waiver of NNI's general or specific objections.

OBJECTIONS AND RESPONSES TO INDIVIDUAL INTERROGATORIES <u>Interrogatory No. 4:</u>

Identify the names and state the present and/or last known address(es) of the individual(s)/employee(s) with the most knowledge pertaining to VICTOZA, including but not limited to:

- (a) The Product managers at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (b) The sales representatives (whether nationally, regionally, etc.) at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
 - (i) If the sales representative was a regional position, please identify all regions that Defendant utilized and the person(s) most knowledgeable for each specific region, identifying the individuals by time period; and
 - (ii) Describe the sales and marketing organizational structure utilized by YOU regarding VICTOZA;
- (c) The safety and compliance individuals in charge of reporting ADVERSE EVENTS and complaints of side effects to the FDA or any other agency, and investigating all ADVERSE EVENTS and complaints of side effects at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (d) The person or persons at all times responsible for Quality Assurance with regard to VICTOZA;
- (e) Defendant's liaison(s) to the FDA, whether or not part of the regulatory affairs department, with regard to VICTOZA at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (f) Defendant's researcher(s) and developer(s) responsible for VICTOZA at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;

- (g) Defendant's scientific researcher(s) of VICTOZA at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (h) The person or persons responsible for Defendant's marketing and/or detailing of VICTOZA at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (i) Defendant's Chief Medical Officer at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (j) Defendant's Chief Executive Officer ("CEO") at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (k) Defendant's President at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (1) Defendant's Chief Financial Officer ("CFO") at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (m) Defendant's Chief Information Officer ("CIO") at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (n) The person responsible for regulatory affairs at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (o) Defendant's liaison(s) with any subsidiary or affiliate located outside the United States that manufactured, produced, promoted, formulated, created,

- designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (p) Defendant's General Counsel and/or the names of all associate general counsel at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period;
- (q) Defendant's Chief Operating Officer ("COO") at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period; and
- (r) Members of any International Product Team maintained or utilized by YOU at all times Defendant manufactured, produced, promoted, formulated, created, designed, sold and/or tested VICTOZA, identifying the individuals by time period.

Response to Interrogatory No. 4:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI further objects to this interrogatory as vague and ambiguous as it fails to define certain terms and phrases, including "complaints of side effects," "liaison," "responsible," and "detailing." NNI further objects to this interrogatory to the extent that it seeks information about Victoza's® manufacture as it is neither relevant nor reasonably calculated to lead to the discovery of admissible evidence because Plaintiffs did not allege any manufacturing defect of Victoza® in this litigation. NNI further objects to the extent this interrogatory seeks information concerning activities outside the United States. NNI objects further to this interrogatory to the extent that it seeks information. NNI objects further to this interrogatory to the extent that it seeks information about Victoza® sales representatives more appropriate for case-specific discovery. NNI objects further to this interrogatory to the extent that it seeks information about Victoza's® International Product Team.

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Subject to and without waiving or otherwise limiting the foregoing general and specific objections, NNI refers Plaintiffs to the organizational charts that will be produced by NNI in this litigation. In addition, pursuant to the parties' agreement on February 3, 2014, NNI will answer this interrogatory more fully at a later time, as needed, once its discovery and document production are more substantially completed and upon mutual agreement by the parties regarding, among other things, the appropriate scope and context of this request.

Supplemental Response to Interrogatory No. 4:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI further objects to this interrogatory as vague and ambiguous as it fails to define certain terms and phrases, including "complaints of side effects," "liaison," "responsible," and "detailing." NNI further objects to this interrogatory to the extent that it seeks information about Victoza's® manufacture as it is neither relevant nor reasonably calculated to lead to the discovery of admissible evidence because Plaintiffs did not allege any manufacturing defect of Victoza® in this litigation. NNI further objects to the extent this interrogatory seeks information concerning activities outside the United States. NNI objects further to this interrogatory to the extent it requests public information. NNI objects further to this interrogatory to the extent that it seeks information about Victoza® sales representatives more appropriate for case-specific discovery. NNI objects further to this interrogatory to the extent that it seeks information about Victoza's® International Product Team. NNI objects further to this interrogatory to the extent it is outside the scope of general causation discovery and inapplicable discovery at this time pursuant to this Court's February 18 Order, which requires Plaintiffs to "narrow all discovery related requests to issues involving general causation." NNI will supplement its answers with information beyond general causation at the appropriate time, if necessary and applicable, per further Order of this Court.

Subject to and without waiving or otherwise limiting the foregoing general

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and specific objections, NNI states that it has numerous employees who may have knowledge relating to Victoza® as addressed in this interrogatory, and it is not possible to identify every such person, particularly at this stage of the litigation. NNI will be producing documents in response to various Requests for Production and as stated in its counsel's February 21, 2014 letter, including documents from custodial files as well as organization charts, and such documents will include the names of many such people from where further answers may be derived. NNI identifies the following individuals as those most likely to have primary knowledge relating to Victoza® with respect to the subject matters identified below:

Regulatory Affairs

 Michelle Thompson: Senior Director, Regulatory Affairs 800 Scudders Mill Road

Plainsboro, NJ 08536

Medical Affairs

 Jason H. Brett, M.D.: Senior Medical Director, Medical Affairs 800 Scudders Mill Road

Plainsboro, NJ 08536

Product Safety

• Michael Sacco: Senior Director, Product Safety

800 Scudders Mill Road

Plainsboro, NJ 08536

Clinical Development

• Christian Born Djurhuus, M.D., Ph.D.: Executive Director, Clinical Development-Diabetes

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- Camille Lee: Senior Vice President, Diabetes Marketing;
- Curt Oltmans: Corporate Vice President and General Counsel;
- Anne Phillips, M.D.: Senior Vice President, Clinical Medical & Regulatory Affairs;
- Eddie Williams: Senior Vice President, Biopharmaceuticals.

Interrogatory No. 5:

Identify all persons and/or entities paid by YOU for consulting services of any kind concerning VICTOZA, and for each such person or entity state the nature of the consulting services rendered and the time frame(s) during which they were rendered.

Response to Interrogatory No. 5:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI objects further to this interrogatory because the term "consulting services of any kind" is overbroad, ambiguous and vague.

Supplemental Response to Interrogatory No. 5:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI objects further to this interrogatory because the term "consulting services of any kind" is overbroad, ambiguous and vague. NNI objects further to this interrogatory to the extent it is outside the scope of general causation discovery and inapplicable discovery at this time pursuant to this Court's February 18 Order, which requires Plaintiffs to "narrow all discovery related requests to issues involving general causation." NNI will supplement its answers with information beyond general causation at the appropriate time, if necessary and applicable, per further Order of this Court. NNI objects further to this interrogatory to the extent that it requests the identification of consultants relevant to individual plaintiffs' treating physicians because that information will be produced as part of the Defense Fact Sheet at the appropriate time per agreement of the parties and Order of this Court.

Subject to and without waiving the foregoing general and specific objections, NNI states that it consults with a range of individuals and entities paid for various services concerning Victoza® including Key Opinion Leaders and Investigators whom NNI will identify in the future, to the extent this information is reasonably accessible. At this time, NNI's investigation into this area is still ongoing and NNI plans to supplement with responsive information relevant for general causation pursuant to the discovery schedule set by this Court on March 4, 2014, on a rolling basis, to the extent information is identified. *See* Order Following Second Case Management Conference, *In re Incretin-Based Therapies Prods. Liab. Litig.*, No. 13-md-2452-AJB-MDD (S.D. Cal. March 4, 2013), Doc. No. 340 ("March 4 Order").

Interrogatory No. 10:

Identify each and every database that YOU or others acting on YOUR behalf maintain or have maintained that is likely to contain any data or information about BYETTA, JANUVIA, JANUMET, VICTOZA and/or any other GLP-1 agonist or DPP-4 inhibitor. Include in your answer:

- (a) The name of each database;
- (b) The identity of the database administrators;
- (c) The dates of use for each database;
- (d) The hardware and software platforms each database utilized;
- (e) The type of information about BYETTA, JANUVIA, JANUMET, VICTOZA, and/or any other GLP-1 agonist or DPP-4 inhibitor contained in each database;
- (f) Whether each database was a transactional database;
- (g) Whether each database was a warehouse database;
- (h) The identity of all other databases that fed information into each database identified;
- (i) The search capabilities of each database;
- (j) The back-up schedule for each database;
- (k) Whether each database has an audit trail feature that has been enabled;

- (1) The archival, retention and destruction policies with respect to each database; and,
- (m) Whether any database has been discontinued and what was done with the data contained in any retired database.

Response to Interrogatory No. 10:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI objects to this interrogatory because it is overly broad and unduly burdensome. NNI further objects to this interrogatory as vague and ambiguous as it fails to define certain terms and phrases, including "transactional database," and "warehouse database." NNI further objects to the extent that this interrogatory relates to or seeks information regarding products other than Victoza®.

Subject to and without waiving or otherwise limiting the foregoing general and specific objections, and pursuant to the parties' agreement on February 3, 2014, NNI will answer this interrogatory more fully at a later time, as needed, once its discovery and document production are more substantially completed and upon mutual agreement by the parties regarding, among other things, the appropriate scope and context of this request.

Supplemental Response to Interrogatory No. 10:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI objects to this interrogatory because it is overly broad and unduly burdensome. NNI further objects to this interrogatory as vague and ambiguous as it fails to define certain terms and phrases, including "transactional database," and "warehouse database." NNI further objects to the extent that this interrogatory relates to or seeks information regarding products other than Victoza®. NNI objects further to this interrogatory to the extent it is outside the scope of general causation discovery and inapplicable discovery at this time pursuant to this Court's February 18 Order which requires Plaintiffs to "narrow all discovery related requests to issues involving general causation." NNI will supplement its answers with

information beyond general causation at the appropriate time, if necessary and applicable, per further Order of this Court.

Subject to and without waiving or otherwise limiting the foregoing general and specific objections, NNI states that it is continuing to investigate and identify the databases where information about Victoza® may be contained and will supplement this response as additional information about NNI's databases is discovered. NNI further states at this time that data or information about Victoza® may be contained in the following databases currently used by NNI:

- ARGUS: contains Victoza® adverse event data;
- IRMS: contains Physician Information Requests concerning Victoza®;
- Insight Manager: contains Regulatory submissions and communications with the FDA regarding Victoza®;
- NovoDocs: contains clinical protocols and clinical trial reports, among other information;
- QBIC: contains NNI's SOPs;
- KOL database: maintains information on KOLs regarding Victoza®;
- CCGLOW: tracks information regarding customer complaints regarding Victoza®; and
- IIS: tracks information regarding Investigator Initiated Studies involving Victoza®.

Interrogatory No. 23:

Did the FDA or any advisory committee or sub-committee of the FDA or any other governmental body ever hold any hearings as to the safety and/or efficacy of BYETTA, JANUVIA, JANUMET and/or VICTOZA? If yes, identify the date(s), time(s), place(s), and participants in the hearings; state whether YOU or anyone acting on YOUR behalf provided testimony at any such hearings (including but not limited to hearings by the FDA, CDC, NIH, USDA, U.S. Congress, and/or U.S. Senate); state the outcome of the hearings; attach all transcripts of such hearings in EASTIVE 224334.4

NOVO NORDISK INC.'S SUPPL. OBJ. AND RESP. TO PLS' AMENDED SECOND SET OF INTERROGATORIES

native electronic form; and state whether the FDA and/or any other governmental body ever suggested, requested, or required YOU to provide further information and/or perform further tests as to the safety of BYETTA, JANUVIA, JANUMET and/or VICTOZA.

Response to Interrogatory No. 23:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI further objects to this interrogatory as vague and ambiguous as it fails to define certain terms, such as "advisory board." NNI further objects to the extent this interrogatory calls for information either not within NNI's possession, custody, or control and/or information that is a matter of public record or otherwise as accessible to Plaintiffs as to NNI. NNI objects further to this interrogatory to the extent it seeks information about regulatory processes and activities outside of the United States because it is neither relevant nor reasonably calculated to lead to the discovery of admissible evidence, and is unduly burdensome to produce in this litigation. NNI further objects to the extent that this interrogatory relates to or seeks information regarding products other than Victoza®.

Subject to and without waiving or otherwise limiting the foregoing general and specific objections, and pursuant to the parties' agreement on February 3, 2014, NNI will answer this interrogatory more fully at a later time, as needed, once its discovery and document production are more substantially completed and upon mutual agreement by the parties regarding, among other things, the appropriate scope and context of this request.

Supplemental Response to Interrogatory No. 23:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI further objects to this interrogatory as vague and ambiguous as it fails to define certain terms, such as "hearings" and "advisory board." NNI further objects to the extent this interrogatory calls for information either not within NNI's possession, custody, or control and/or information that is a matter of public record or

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otherwise as accessible to Plaintiffs as to NNI. NNI objects further to this interrogatory to the extent it seeks information about regulatory processes and activities outside of the United States because it is neither relevant nor reasonably calculated to lead to the discovery of admissible evidence, and is unduly burdensome to produce in this litigation. NNI further objects to investigating this interrogatory to the extent it is intended to refer to FDA and government hearings which are matters of public record that are as accessible to plaintiffs as to NNI. NNI further objects to the extent that this interrogatory relates to or seeks information regarding products other than Victoza®. NNI objects further to this interrogatory to the extent it is outside the scope of general causation discovery and inapplicable discovery at this time pursuant to this Court's February 18 Order, which requires Plaintiffs to "narrow all discovery related requests to issues involving general causation." NNI will supplement its answers with information beyond general causation at the appropriate time, if necessary and applicable, per further Order of this Court.

Subject to and without waiving or otherwise limiting the foregoing general and specific objections, NNI states that on June 12-13, the National Institute of Diabetes and Digestive and Kidney Diseases (NIDDK) and the National Cancer Institute cosponsored a workshop at the National Institutes of Health (the "Workshop"). The purpose of the Workshop was to bring together experts from academia, medicine, industry, and governmental agencies to present relevant data and perspectives on questions related to diabetes and the risk of pancreatic cancer. The Workshop included a session on the effects of incretin-based therapies. Global Chief Medical Officer Alan C. Moses, M.D. attended the workshop and spoke on behalf of NNI during this session in a presentation titled, "PDAC Incidence with Liraglutide Therapy." See Session 6: Effects of DM Treatment on PDAC, NIDDK-NCI Workshop on Pancreatitis-Diabetes-Pancreatic Cancer, NIDDK, available at http://www.niddk.nih.gov/news/events-calendar/Pages/niddknci-workshop-on-

additional employees may have attended the workshop but did not participate or make a presentation. Their names are included in the materials provided to all attendees, of which we understand Plaintiffs are in possession.

Interrogatory No. 39:

Identify each person acting on YOUR behalf who has been responsible for: (a) receiving any complaints, inquiries, letters and other documents pertaining to VICTOZA; (b) evaluating any complaints, inquiries, letters, and other documents pertaining to VICTOZA; (c) investigating any complaints, inquiries, letters or other documents pertaining to VICTOZA; and (d) responding to any complaints, inquiries, letters and other documents pertaining to VICTOZA.

Response to Interrogatory No. 39:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI further objects to this interrogatory as vague and ambiguous as it fails to define certain terms and phrases, including "on your behalf," "responsible," "inquiries," "complaints" and "responding." NNI further objects to this interrogatory to the extent it seeks information unrelated to the alleged risks and injuries at issue in this litigation. NNI further objects to this interrogatory to the extent it seeks information concerning activities outside the United States because it is neither relevant nor reasonably calculated to lead to the discovery of admissible evidence and is unduly burdensome.

Subject to and without waiving or otherwise limiting the foregoing general and specific objections, and pursuant to the parties' agreement on February 3, 2014, NNI will answer this interrogatory more fully at a later time, as needed, once its discovery and document production are more substantially completed and upon mutual agreement by the parties regarding, among other things, the appropriate scope and context of this request.

Supplemental Response to Interrogatory No. 39:

NNI incorporates, as if fully set forth herein, the General Objections by
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reference. NNI further objects to this interrogatory as vague and ambiguous as it fails to define certain terms and phrases, including "on your behalf," "responsible," "inquiries," "complaints" and "responding." NNI further objects to this interrogatory to the extent it seeks information unrelated to the alleged risks and injuries at issue in NNI further objects to this interrogatory to the extent it seeks this litigation. information concerning activities outside the United States because it is neither relevant nor reasonably calculated to lead to the discovery of admissible evidence and is unduly burdensome. NNI objects further to this interrogatory to the extent it is outside the scope of general causation discovery and inapplicable discovery at this time pursuant to this Court's February 18 Order, which requires Plaintiffs to "narrow" all discovery related requests to issues involving general causation." NNI will supplement its answers with information beyond general causation at the appropriate time, if necessary and applicable, per further Order of this Court.

Subject to and without waiving or otherwise limiting the foregoing general and specific objections, NNI responds that various employees and departments within NNI receive, evaluate, investigate and/or respond to complaints, inquiries, and/or letters regarding Victoza®, and that the Product Safety department is ultimately responsible for all complaints regarding Victoza® marketed in the U.S. Michael Sacco, Senior Director of Product Safety, has responsibility for the Product Safety department.

Interrogatory No. 48:

At any time since VICTOZA became publicly available in the United States, have YOU discussed or considered withdrawing it from the market due to reports of ADVERSE EVENTS or for any other reason? If YOUR answer is in the affirmative, please state:

- When withdrawal was discussed or considered; (a)
- (b) Who was involved in any discussions regarding withdrawal;
- What prompted any discussions regarding withdrawal;

- (d) Whether any studies were undertaken or reviewed in discussing or considering withdrawal and, if so, identify which ones; and
- (e) Why it was determined not to withdraw VICTOZA from the United States market.

Response to Interrogatory No. 48:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI objects to the extent this interrogatory requests information concerning activities outside the United States. NNI further objects to this interrogatory to the extent it is unduly burdensome.

Subject to and without waiving or otherwise limiting the foregoing general and specific objections, and pursuant to the parties' agreement on February 3, 2014, NNI will answer this interrogatory more fully at a later time, as needed, once its discovery and document production are more substantially completed and upon mutual agreement by the parties regarding, among other things, the appropriate scope and context of this request.

Supplemental Response to Interrogatory No. 48:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI objects to the extent this interrogatory requests information concerning activities outside the United States. NNI further objects to this interrogatory to the extent it is unduly burdensome. NNI objects further to this interrogatory to the extent it is outside the scope of general causation discovery and inapplicable discovery at this time pursuant to this Court's February 18 Order, which requires Plaintiffs to "narrow all discovery related requests to issues involving general causation." NNI will supplement its answers with information beyond general causation at the appropriate time, if necessary and applicable, per further Order of this Court.

Subject to and without waiving or otherwise limiting the foregoing general and specific objections, NNI states that it monitors and evaluates the safety data

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28 (b) When the removal(s) occurred;

concerning Victoza® on an ongoing basis and to date, has determined that Victoza® does not need to be withdrawn from the market due to safety concerns or otherwise.

NNI further states that its decisions are consistent with the July 25, 2013 EMA Assessment Report, which states that the pre-clinical data "do not indicate that [Victoza] treatment is associated with adverse effects on the [] pancreas" and that "there is currently no support from clinical trials that [incretin] based therapies increase the risk" of pancreatic cancer. See European Medicines Agency, Assessment Report for GLP-1 Based Therapies, July 25, 2013, at 8, 16. NNI further states that its decisions are consistent with the recent conclusions from the FDA and EMA published on February 27, 2014 in the New England Journal of Medicine, which state that "the FDA and the EMA have explored multiple streams of data pertaining to a pancreatic safety signal associated with incretin-based drugs. Both agencies agree that assertions concerning a causal association between incretin-based drugs and pancreatitis or pancreatic cancer, as expressed recently in the scientific literature and in the media, are inconsistent with the current data.... The FDA and the EMA have not reached a final conclusion at this time regarding such a causal relationship... The FDA and the EMA believe that the current knowledge is adequately reflected in the product information or labeling." See Amy Egan, M.D., M.P.H., et al. Pancreatic Safety of Incretin-Based Drugs –FDA and EMA Assessment, 370:9 New. Eng. J. Med. 794, 796 (2014). NNI further refers Plaintiffs to its NDA and IND productions which include communications with the FDA regarding the safety of Victoza®.

Interrogatory No. 49:

Has there ever been a discontinuance, either temporary or otherwise, of any exenatide, sitagliptin, liraglutide and/or any other GLP-1 agonist or DPP-4 inhibitor-containing medication in the United States or any other country? If YOUR answer is in the affirmative, indicate the following:

- (a) Which drug(s) were removed from the market;
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- (c) Whether the discontinuance(s) were permanent or temporary;
- (d) The primary motivations behind the discontinuance(s); and,
- (e) The rate of discontinuance in comparison to the overall prevalence of the drug(s) on the market.

Response to Interrogatory No. 49:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI objects to this interrogatory as vague and ambiguous. NNI further objects to this interrogatory to the extent it seeks information unrelated to the alleged risks and injuries at issue in this litigation. NNI further objects to this interrogatory to the extent it seeks information concerning products other than Victoza®. NNI further objects to the extent this interrogatory requests information regarding foreign regulatory activities.

Subject to and without waiving or otherwise limiting the foregoing general and specific objections, and pursuant to the parties' agreement on February 3, 2014, NNI will answer this interrogatory more fully at a later time, as needed, once its discovery and document production are more substantially completed and upon mutual agreement by the parties regarding, among other things, the appropriate scope and context of this request.

Supplemental Response to Interrogatory No. 49:

NNI incorporates, as if fully set forth herein, the General Objections by reference. NNI objects to this interrogatory as vague and ambiguous. NNI further objects to this interrogatory to the extent it seeks information unrelated to the alleged risks and injuries at issue in this litigation. NNI further objects to this interrogatory to the extent it seeks information concerning products other than Victoza®. NNI further objects to the extent this interrogatory requests information regarding foreign regulatory activities. NNI objects further to this interrogatory to the extent it is outside the scope of general causation discovery and inapplicable discovery at this time pursuant to this Court's February 18 Order, which requires Plaintiffs to "narrow

all discovery related requests to issues involving general causation." NNI will supplement its answers with information beyond general causation at the appropriate time, if necessary and applicable, per further Order of this Court.

Subject to and without waiving or otherwise limiting the foregoing general and specific objections, NNI states that there has not been any discontinuance of Victoza® in the U.S. market.

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2	Dated: March 6, 2014	DLA PIPER LLP (US)	
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4		By: /s/ Raymond M. Williams	
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		(Bar No. 164068)	
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25		Nordisk Inc.	U
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CERTIFICATE OF SERVICE

2 I, Lauren M. Nonnemacher, declare:

I am a citizen of the United States and employed in the county of Philadelphia, state of Pennsylvania. I am over the age of eighteen years and not a party to the within-entitled action. My business address is DLA Piper LLP (US), One Liberty Place, 1650 Market Street, Suite 4900, Philadelphia, PA 19103. On March 6, 2014, I served a copy of the within document:

Defendant Novo Nordisk Inc.'s Supplemental Objections and Responses to Plaintiffs' Amended Second Set of Interrogatories

- by placing the document listed above in a sealed envelope with postage thereon fully prepaid, the United States mail at Philadelphia, Pennsylvania addressed as set forth below.
- by placing the document listed above in a sealed Delivery Service envelope and affixing a pre-paid air bill, and causing the envelope to be delivered to a Delivery Service agent for delivery.
- by personally delivering the document listed above to the persons at the address set forth below.
- I hereby certify that on the below date, I electronically filed the foregoing with the Clerk of the Court using the CM/ECF system which will send notification of such filing to the e-mail addresses denoted on the Electronic Mail notice list, and I hereby certify that I have mailed the foregoing document or paper via the United States Postal Service to the non-CM/ECF participants indicated on the Manual Notice list (if any).
- by transmitting via e-mail or electronic transmission the document listed above to the persons at the e-mail addresses set forth below.

1 2	Gayle M. Blatt CASEY GERRY SCHENK FRANCAVILLA BLATT &	Michael K. Johnson JOHNSON BECKER, PLLC 33 South Sixth Street, Suite 4530
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4	San Diego, CA 92101 gmb@cglaw.com Plaintiffs' Co-Liaison Counsel	Tiumigs Executive Committee
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9	Hunter J. Shkolnik NAPOLI BERN RIPKA SHKOLNIK	Amy J. Laurendeau Scott M. Edson
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17	kingk@pepperlaw.com Attorneys for Defendant Eli Lilly and	Washington, DC 20005 dmarvin@wc.com
18	Company Company	eesber@wc.com pboehm@wc.com
19		Attorneys for Defendant Merck Sharp & Dohme Corp.
20		
21	I am readily familiar with the firm's practice of collection and processin	
22	correspondence for mailing. Under that practice it would be deposited with the	
23	U.S. Postal Service on that same day	with postage thereon fully prepaid in the
24	ordinary course of business. I am awar	re that on motion of the party served, service
25	is presumed invalid if postal cancellat	ion date or postage meter date is more than
26	one day after date of deposit for mailing in affidavit.	
27	I declare that I am employed in the office of a member of the bar of this court	

at whose direction the service was made.

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1	Executed on March 6, 2014, at Philadelphia, Pennsylvania.
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4	/s/ Lauren M. Nonnemacher Lauren M. Nonnemacher
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	-3- DEFENDANT NOVO NORDISK INC.'S SUPPLEMENTAL OBJECTIONS AND RESPONSES TO